

# Aleksandra Trifunovic

## List of Publications by Year in descending order

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Version: 2024-02-01

70  
papers

8,423  
citations

145106

33  
h-index

107981

68  
g-index

76  
all docs

76  
docs citations

76  
times ranked

12278  
citing authors

#	ARTICLE	IF	CITATIONS
1	CLPP deficiency ameliorates neurodegeneration caused by impaired mitochondrial protein synthesis. <i>Brain</i> , 2022, 145, 92-104.	3.7	9
2	Mitochondrial matrix proteases: quality control and beyond. <i>FEBS Journal</i> , 2022, 289, 7128-7146.	2.2	27
3	Spatial and temporal control of mitochondrial H <sub>2</sub> O <sub>2</sub> release in intact human cells. <i>EMBO Journal</i> , 2022, 41, e109169.	3.5	39
4	FGF21 modulates mitochondrial stress response in cardiomyocytes only under mild mitochondrial dysfunction. <i>Science Advances</i> , 2022, 8, eabn7105.	4.7	23
5	CLUH controls astrin-1 expression to couple mitochondrial metabolism to cell cycle progression. <i>ELife</i> , 2022, 11, .	2.8	7
6	Remission of obesity and insulin resistance is not sufficient to restore mitochondrial homeostasis in visceral adipose tissue. <i>Redox Biology</i> , 2022, 54, 102353.	3.9	14
7	Host-commensal interaction promotes health and lifespan in <i>Caenorhabditis elegans</i> through the activation of HLH-30/TFEB-mediated autophagy. <i>Aging</i> , 2021, 13, 8040-8054.	1.4	15
8	Tune instead of destroy: How proteolysis keeps OXPHOS in shape. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2021, 1862, 148365.	0.5	29
9	DNA polymerase gamma mutations that impair holoenzyme stability cause catalytic subunit depletion. <i>Nucleic Acids Research</i> , 2021, 49, 5230-5248.	6.5	15
10	Phosphoproteomics of the developing heart identifies PERM1 - An outer mitochondrial membrane protein. <i>Journal of Molecular and Cellular Cardiology</i> , 2021, 154, 41-59.	0.9	9
11	Adaptation to mitochondrial stress requires CHOP-directed tuning of ISR. <i>Science Advances</i> , 2021, 7, .	4.7	68
12	Systemic regulation of mitochondria by germline proteostasis prevents protein aggregation in the soma of <i>C. elegans</i> . <i>Science Advances</i> , 2021, 7, .	4.7	28
13	PERM1 interacts with the MICOS-MIB complex to connect the mitochondria and sarcolemma via ankyrin B. <i>Nature Communications</i> , 2021, 12, 4900.	5.8	6
14	Implication of folate deficiency in CYP2U1 loss of function. <i>Journal of Experimental Medicine</i> , 2021, 218, .	4.2	13
15	Adaptive translational pausing is a hallmark of the cellular response to severe environmental stress. <i>Molecular Cell</i> , 2021, 81, 4191-4208.e8.	4.5	18
16	Mitochondrial metabolism coordinates stage-specific repair processes in macrophages during wound healing. <i>Cell Metabolism</i> , 2021, 33, 2398-2414.e9.	7.2	89
17	Identification of Putative Mitochondrial Protease Substrates. <i>Methods in Molecular Biology</i> , 2021, 2192, 313-329.	0.4	1
18	Neuronal ablation of mt-AspRS in mice induces immune pathway activation prior to severe and progressive cortical and behavioral disruption. <i>Experimental Neurology</i> , 2020, 326, 113164.	2.0	11

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19	A <i>C. elegans</i> model for neurodegeneration in Cockayne syndrome. <i>Nucleic Acids Research</i> , 2020, 48, 10973-10985.	6.5	23
20	DARS2 is indispensable for Purkinje cell survival and protects against cerebellar ataxia. <i>Human Molecular Genetics</i> , 2020, 29, 2845-2854.	1.4	11
21	Mitochondrial DNA mutations and aging. , 2020, , 221-242.		5
22	Mitochondrial Regulation of the 26S Proteasome. <i>Cell Reports</i> , 2020, 32, 108059.	2.9	28
23	GLP-1 Receptor Signaling in Astrocytes Regulates Fatty Acid Oxidation, Mitochondrial Integrity, and Function. <i>Cell Metabolism</i> , 2020, 31, 1189-1205.e13.	7.2	76
24	The Mouse Heart Mitochondria N Terminome Provides Insights into ClpXP-Mediated Proteolysis. <i>Molecular and Cellular Proteomics</i> , 2020, 19, 1330-1345.	2.5	20
25	Mitochondrial Dysfunction Combined with High Calcium Load Leads to Impaired Antioxidant Defense Underlying the Selective Loss of Nigral Dopaminergic Neurons. <i>Journal of Neuroscience</i> , 2020, 40, 1975-1986.	1.7	34
26	A salvage pathway maintains highly functional respiratory complex I. <i>Nature Communications</i> , 2020, 11, 1643.	5.8	80
27	Alterations of redox and iron metabolism accompany the development of HIV latency. <i>EMBO Journal</i> , 2020, 39, e102209.	3.5	37
28	Prostaglandin signals from adult germline stem cells delay somatic ageing of <i>Caenorhabditis elegans</i> . <i>Nature Metabolism</i> , 2019, 1, 790-810.	5.1	30
29	KLF-1 orchestrates a xenobiotic detoxification program essential for longevity of mitochondrial mutants. <i>Nature Communications</i> , 2019, 10, 3323.	5.8	25
30	Loss of genomic integrity induced by lysosphingolipid imbalance drives ageing in the heart. <i>EMBO Reports</i> , 2019, 20, .	2.0	26
31	CLPP deficiency protects against metabolic syndrome but hinders adaptive thermogenesis. <i>EMBO Reports</i> , 2018, 19, .	2.0	42
32	Mechanisms of Mitochondria Assembly, Dynamics and Turnover in Health and Disease. <i>Journal of Molecular Biology</i> , 2018, 430, 4821-4822.	2.0	0
33	DARS2 protects against neuroinflammation and apoptotic neuronal loss, but is dispensable for myelin producing cells. <i>Human Molecular Genetics</i> , 2017, 26, 4181-4189.	1.4	23
34	Deficiency of <i>WARS2</i> , encoding mitochondrial tryptophanyl tRNA synthetase, causes severe infantile onset leukoencephalopathy. <i>American Journal of Medical Genetics, Part A</i> , 2017, 173, 2505-2510.	0.7	36
35	Origins of mtDNA mutations in ageing. <i>Essays in Biochemistry</i> , 2017, 61, 325-337.	2.1	54
36	Loss of CLPP alleviates mitochondrial cardiomyopathy without affecting the mammalian UPR <sup>mt</sup> . <i>EMBO Reports</i> , 2016, 17, 953-964.	2.0	86

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37	The CoQH2/CoQ Ratio Serves as a Sensor of Respiratory Chain Efficiency. <i>Cell Reports</i> , 2016, 15, 197-209.	2.9	215
38	<scp>CLPP</scp> coordinates mitoribosomal assembly through the regulation of <scp>ERAL</scp> 1 levels. <i>EMBO Journal</i> , 2016, 35, 2566-2583.	3.5	123
39	Human R1441C LRRK2 regulates the synaptic vesicle proteome and phosphoproteome in a <i>Drosophila</i> model of Parkinson's disease. <i>Human Molecular Genetics</i> , 2016, 25, ddw352.	1.4	61
40	Different faces of mitochondrial DNA mutators. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2015, 1847, 1362-1372.	0.5	39
41	Spastin Binds to Lipid Droplets and Affects Lipid Metabolism. <i>PLoS Genetics</i> , 2015, 11, e1005149.	1.5	84
42	Is mitochondrial free radical theory of aging getting old?. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2015, 1847, 1345-1346.	0.5	9
43	Loss of UCP2 Attenuates Mitochondrial Dysfunction without Altering ROS Production and Uncoupling Activity. <i>PLoS Genetics</i> , 2014, 10, e1004385.	1.5	63
44	Embelin inhibits endothelial mitochondrial respiration and impairs neoangiogenesis during tumor growth and wound healing. <i>EMBO Molecular Medicine</i> , 2014, 6, 624-639.	3.3	71
45	Defects in $\beta$ -Cell Ca <sup>2+</sup> Dynamics in Age-Induced Diabetes. <i>Diabetes</i> , 2014, 63, 4100-4114.	0.3	35
46	Obesity-Induced CerS6-Dependent C16:0 Ceramide Production Promotes Weight Gain and Glucose Intolerance. <i>Cell Metabolism</i> , 2014, 20, 678-686.	7.2	520
47	Tissue-Specific Loss of DARS2 Activates Stress Responses Independently of Respiratory Chain Deficiency in the Heart. <i>Cell Metabolism</i> , 2014, 19, 458-469.	7.2	185
48	Succinate Dehydrogenase Upregulation Destabilize Complex I and Limits the Lifespan of gas-1 Mutant. <i>PLoS ONE</i> , 2013, 8, e59493.	1.1	31
49	Somatic Progenitor Cell Vulnerability to Mitochondrial DNA Mutagenesis Underlies Progeroid Phenotypes in Polg Mutator Mice. <i>Cell Metabolism</i> , 2012, 15, 100-109.	7.2	213
50	Random mtDNA mutations modulate proliferation capacity in mouse embryonic fibroblasts. <i>Biochemical and Biophysical Research Communications</i> , 2011, 409, 394-399.	1.0	13
51	A Tissue-Specific Approach to the Analysis of Metabolic Changes in <i>Caenorhabditis elegans</i> . <i>PLoS ONE</i> , 2011, 6, e28417.	1.1	15
52	High brain lactate is a hallmark of aging and caused by a shift in the lactate dehydrogenase A/B ratio. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2010, 107, 20087-20092.	3.3	218
53	Point Mutations Are Causing Progeroid Phenotypes in the mtDNA Mutator Mouse. <i>Cell Metabolism</i> , 2010, 11, 1.	7.2	12
54	Response: Point Mutations Are Causing Progeroid Phenotypes in the mtDNA Mutator Mouse. <i>Cell Metabolism</i> , 2010, 11, 93.	7.2	5

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55	Caenorhabditis elegans as a model system for mtDNA replication defects. <i>Methods</i> , 2010, 51, 437-443.	1.9	28
56	Mitochondrial DNA level, but not active replicase, is essential for Caenorhabditis elegans development. <i>Nucleic Acids Research</i> , 2009, 37, 1817-1828.	6.5	100
57	Somatic mtDNA mutations and aging " Facts and fancies. <i>Experimental Gerontology</i> , 2009, 44, 101-105.	1.2	33
58	Random Point Mutations with Major Effects on Protein-Coding Genes Are the Driving Force behind Premature Aging in mtDNA Mutator Mice. <i>Cell Metabolism</i> , 2009, 10, 131-138.	7.2	200
59	The mtDNA mutator mouse: Dissecting mitochondrial involvement in aging. <i>Aging</i> , 2009, 1, 1028-1032.	1.4	52
60	Strong Purifying Selection in Transmission of Mammalian Mitochondrial DNA. <i>PLoS Biology</i> , 2008, 6, e10.	2.6	425
61	Progressive parkinsonism in mice with respiratory-chain-deficient dopamine neurons. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2007, 104, 1325-1330.	3.3	516
62	Somatic mtDNA mutations cause progressive hearing loss in the mouse. <i>Experimental Cell Research</i> , 2007, 313, 3924-3934.	1.2	48
63	Mitochondrial DNA and ageing. <i>Biochimica Et Biophysica Acta - Bioenergetics</i> , 2006, 1757, 611-617.	0.5	96
64	Proteolytic Processing of OPA1 Links Mitochondrial Dysfunction to Alterations in Mitochondrial Morphology. <i>Journal of Biological Chemistry</i> , 2006, 281, 37972-37979.	1.6	382
65	Mitochondrial DNA polymerase gamma is essential for mammalian embryogenesis. <i>Human Molecular Genetics</i> , 2005, 14, 1775-1783.	1.4	219
66	Modelling the mitochondrial role in ageing. <i>Drug Discovery Today: Disease Models</i> , 2005, 2, 257-263.	1.2	0
67	Somatic mtDNA mutations cause aging phenotypes without affecting reactive oxygen species production. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2005, 102, 17993-17998.	3.3	491
68	Premature ageing in mice expressing defective mitochondrial DNA polymerase. <i>Nature</i> , 2004, 429, 417-423.	13.7	2,318
69	Tissue-Specific Knockout Model for Study of Mitochondrial DNA Mutation Disorders. <i>Methods in Enzymology</i> , 2002, 353, 409-421.	0.4	9
70	Mitochondrial transcription factors B1 and B2 activate transcription of human mtDNA. <i>Nature Genetics</i> , 2002, 31, 289-294.	9.4	535